

Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

Claim 1 (Cancelled).

2 (Currently amended). ~~Method of preparation of A~~
method for preparing a population of circulating CD34⁺ cells
~~capable of regenerating which regenerate~~ hematopoiesis in vivo,
comprising:

a) administering to a donor a composition comprising growth hormone or one of its derivatives or any factor inducing growth hormone release, simultaneously or separately with a composition comprising G-CSF, in an amount sufficient to enhance the mobilization or peripheralization effect of G-CSF to further increase in said donor the number of circulating CD34⁺ cells
~~capable of regenerating which regenerate~~ hematopoiesis in vivo[[.]] beyond that achieved by G-CSF alone; and

b) isolating a population of circulating CD34⁺ cells
~~capable of regenerating which regenerate~~ hematopoiesis in vivo
from the peripheral blood of said donor.

Claims 3 and 4 (Cancelled).

5 (Currently amended). ~~Method~~ The method according to claim [[4]] 2, wherein the increased number of CD34⁺ cells is more than 10, 25, 34 or 80 CD34⁺ cells per microliter of donor peripheral blood.

6 (Currently amended). ~~Method~~ The method according to claim [[4]] 2, wherein the increased number of CD34⁺ cells is at least 2×10^6 , 4×10^6 , 5×10^6 , 6×10^6 , 8×10^6 , 15×10^6 CD34⁺ cells per kilogram of donor body weight.

7 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the increased number of circulating CD34⁺ cells ~~capable of regenerating~~ which regenerate hematopoiesis *in vivo* corresponds to ~~around or more than~~ about 500 or more CFU-GM per milliliter of donor peripheral blood.

8 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the increased number of circulating CD34⁺ cells ~~capable of regenerating~~ which regenerate hematopoiesis *in vivo* corresponds to an increased level of CFU-C, CFU-Meg or BFU-E in donor peripheral blood.

9 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the increased number of circulating CD34⁺ cells ~~capable of regenerating~~ which regenerate hematopoiesis *in vivo* substantially corresponds to a white blood

cell count of ~~around or more than~~ about 1000 or more cells per microliter of donor peripheral blood.

10 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the increased number of circulating CD34⁺ cells capable of regenerating which regenerate hematopoiesis *in vivo* corresponds to ~~around or more than~~ about 1×10^5 or more GM-CFC per kilogram of donor or recipient body weight.

11 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the circulating CD34⁺ cells capable of regenerating which regenerate hematopoiesis *in vivo* are CD34⁺/CD33⁺ cells and/or CD34⁺/CD38⁻ cells and/or CD34⁺/Thy-I cells and/or CD34⁺/Thy-I/CD38⁻ cells ~~and/or CD33⁺ cells~~ and/or bone-marrow stem cells and/or progenitor cells and/or long-term culture initiating cells (LTC-IC) and/or cells that fulfill self renewal potential and/or cells that fulfill pluripotential characteristics and/or cells that initiate long term bone marrow culture and/or cells that can generate multiple cell lineages.

12 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the target number of circulating CD34⁺ cells capable of regenerating which regenerate hematopoiesis *in vivo* is at least 2×10^4 LTC-IC per kg of donor or

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recipient body, ~~around or more than~~ about 2×10^6 or more CD34⁺ cells per kilogram of donor or recipient body weight, ~~around or more than~~ about 4×10^6 or more CD34⁺ cells per kilogram of donor or recipient body weight or ~~around or more than~~ about 8×10^6 or more CD34⁺ cells per kilogram of donor or recipient body weight.

13(Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the volume of blood processed in step (b) is ~~emprised~~ in a range of about 30 to about 900 milliliters.

Claims 14-17 (Cancelled).

18(Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein growth-hormone or one of its derivatives or any factor inducing growth hormone release is administered in an amount ~~emprised~~ between 20 to 50 $\mu\text{g/kg}$ of donor body weight, in an amount ~~emprised~~ between 30 to 40 $\mu\text{g/kg}$ of donor body weight or in an amount of 33 μg per kilogram of donor body weight.

19(Currently amended). ~~Method~~ The method according to claim ~~[[17]]~~ 2, wherein the G-CSF is administered in an amount ~~emprised~~ between 3 to 15 $\mu\text{g/kg}$ of donor body weight, in an amount ~~emprised~~ between 4 to 12 $\mu\text{g/kg}$ of donor body weight or in an amount of around 5 or 10 μg per kilogram of donor body weight.

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20 (Currently amended). ~~Method~~ The method according to claim 2, wherein the administration of Growth Hormone is made three times a day and the administration of G-CSF is made daily.

21 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the administration of said composition is made by parenteral, subcutaneous, intravenous, intramuscular, intraperitoneal, transdermal or buccal routes.

22 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ wherein the administration of said composition is daily or three times a day.

23 (Currently amended). ~~Method~~ The method according to ~~claim 1 or 2~~, wherein the administration of said composition is made over a period of 5 days or over a period of 10 days.

24 (Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the administration begins around 7 days after the beginning of a chemotherapeutic treatment or around 2 days after the end of a chemotherapeutic treatment.

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25(Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the growth hormone is a recombinant growth hormone.

26(Currently amended). ~~Method~~ The method according to ~~any one of claims 1 to 3~~ claim 2, wherein the growth hormone is human growth hormone.

Claims 27-55 (Cancelled).

56(New). The method according to claim 2, wherein said growth hormone or one of its derivatives or any factor inducing growth hormone release is administered at a different time than said composition comprising G-CSF.

57(New). The method according to claim 2, wherein said growth hormone or one of its derivatives or any factor inducing growth hormone release is administered simultaneously with said composition comprising G-CSF.